

> d his

(FILE 'HOME' ENTERED AT 07:39:55 ON 20 JAN 2004)

FILE 'MEDLINE, CAPLUS, BIOSIS, EMBASE, SCISEARCH, AGRICOLA'  
ENTERED AT  
07:40:20 ON 20 JAN 2004

L1 555584 S GLYCOPROTEIN  
L2 4677 S ALBUMEN (P) EGG  
L3 17366 S WHEY (P) MILK  
L4 237 S L1 (P) (L2 OR L3)  
L5 98254 S (HELICOBACTER PYLORI) OR (H. PYLORI)  
L6 9072 S UREASE (P) L5  
L7 1 S L4 (P) L6  
L8 3915 S L5 (P) COLONIZATION  
L9 597 S L8 (P) INHIBIT?  
L10 1 S L4 (P) L9  
L11 0 S L10 NOT L7  
L12 40948 S GASTROINTESTINAL DISEASE  
L13 0 S L12 (P) L4  
L14 186 S (HIGH MOLECULAR WEIGHT) (P) WHEY  
L15 24 S (HIGH MOLECULAR WEIGHT) (P) ALBUMEN  
L16 19 S (L14 OR L15) (P) L1  
L17 0 S L16 (P) L6  
L18 3393 S KODAMA Y?/AU  
L19 4890 S KIMURA N?/AU  
L20 1 S (L18 OR L19) AND L4  
L21 0 S L20 NOT L10  
L22 29 S (L18 OR L19) AND L6  
L23 6 S L22 AND L1  
L24 2 DUPLICATE REMOVE L23 (4 DUPLICATES REMOVED)  
L25 1 S L24 NOT L10

=> log y



which specifically binds to \*\*\*H\*\*\* . \*\*\*pylori\*\*\* \*\*\*urease\*\*\*  
 . This \*\*\*glycoprotein\*\*\* is isolated and purified from a  
 \*\*\*glycoprotein\*\*\* -contg. substance, esp. that derived from bovine  
 \*\*\*milk\*\*\* \*\*\*whey\*\*\* or \*\*\*albumen\*\*\* of chicken \*\*\*eggs\*\*\*  
 by affinity chromatog. using a column on which \*\*\*H\*\*\* . \*\*\*pylori\*\*\*  
 \*\*\*urease\*\*\* is immobilized. The \*\*\*glycoprotein\*\*\* is able to  
 effectively inhibit \*\*\*H\*\*\* . \*\*\*pylori\*\*\* colonization, and thus  
 is useful for the prevention or treatment of diseases caused by infection  
 of \*\*\*H\*\*\* . \*\*\*pylori\*\*\* such as peptic ulcers. A food and  
 medicament comprising the inhibitor are also provided.

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=> s 15 (p) colonization

L8 3915 L5 (P) COLONIZATION

=> s 18 (p) inhibit?

L9 597 L8 (P) INHIBIT?

=> s 14 (p) 19

L10 1 L4 (P) L9

=> s 110 not 17

L11 0 L10 NOT L7

=> s gastrointestinal disease

L12 40948 GASTROINTESTINAL DISEASE

=> s 112 (p) 14

L13 0 L12 (P) L4

=> s (high molecular weight) (p) whey

L14 186 (HIGH MOLECULAR WEIGHT) (P) WHEY

=> s (high molecular weight) (p) albumen

L15 24 (HIGH MOLECULAR WEIGHT) (P) ALBUMEN

=> s (114 or 115) (p) 11

L16 19 (L14 OR L15) (P) L1

=> s 116 (p) 16

L17 0 L16 (P) L6

=> s kodama y?/au

L18 3393 KODAMA Y?/AU

=> s kimura n?/au

L19 4890 KIMURA N?/AU

=> s (118 or 119) and 14

L20 1 (L18 OR L19) AND L4

=> s 120 not 110

L21 0 L20 NOT L10

=> s (118 or 119) and 16

L22 29 (L18 OR L19) AND L6

=> s 122 and 11

L23 6 L22 AND L1

=> duplicate remove 123

DUPLICATE PREFERENCE IS 'MEDLINE, CAPLUS, BIOSIS, EMBASE, SCISEARCH'

PROCESSING COMPLETED FOR L23  
L24 2 DUPLICATE REMOVE L23 (4 DUPLICATES REMOVED)

=> s l24 not l10  
L25 1 L24 NOT L10

=> d l25 1 ibib abs

L25 ANSWER 1 OF 1 MEDLINE on STN  
ACCESSION NUMBER: 2000403971 MEDLINE  
DOCUMENT NUMBER: 20389972 PubMed ID: 10930371  
TITLE: Acid-dependent adherence of \*\*\*Helicobacter\*\*\*  
\*\*\*pylori\*\*\* \*\*\*urease\*\*\* to diverse polysaccharides.  
AUTHOR: Icatlo F C; Goshima H; \*\*\*Kimura N\*\*\* ; \*\*\*Kodama Y\*\*\*  
CORPORATE SOURCE: Immunology Research Institute, Ghen Corp., Sano, Gifu City,  
Japan.. irig@ghen.co.jp  
SOURCE: GASTROENTEROLOGY, (2000 Aug) 119 (2) 358-67.  
Journal code: 0374630. ISSN: 0016-5085.  
PUB. COUNTRY: United States  
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)  
LANGUAGE: English  
FILE SEGMENT: Abridged Index Medicus Journals; Priority Journals  
ENTRY MONTH: 200008  
ENTRY DATE: Entered STN: 20000901  
Last Updated on STN: 20000901  
Entered Medline: 20000822

AB BACKGROUND & AIMS: The significance of acid-primed recognition of ligands  
by \*\*\*Helicobacter\*\*\* \*\*\*pylori\*\*\* \*\*\*urease\*\*\* is unknown.  
This study aimed to further characterize the specificity of \*\*\*urease\*\*\*  
adherence in vitro and verify whether specific inhibition will translate  
into in vivo suppression of colonization. METHODS: A highly sensitive  
competitive enzyme-linked ligand capture assay was used to quantify the  
capacity of each test inhibitor to compete with labeled mucin for binding  
sites on immobilized native \*\*\*urease\*\*\*. A model polymer that  
strongly bound \*\*\*urease\*\*\* was used in an in vivo trial using  
euthymic hairless mice as an infection model. RESULTS: The blockage of  
\*\*\*urease\*\*\*-gastric mucin interaction by certain inhibitors revealed an  
acid-functional lectin-like activity by \*\*\*urease\*\*\*, specifically  
recognizing bacterial lipopolysaccharides and certain species of  
polysaccharides, nonbacterial glycolipids, and \*\*\*glycoproteins\*\*\*.  
Dextran sulfate significantly (P < 0.01) suppressed colonization of mice  
by \*\*\*H\*\*\*. \*\*\*pylori\*\*\* when given before and/or after challenge.  
CONCLUSIONS: The acid-driven high-affinity adherence of \*\*\*H\*\*\*  
\*\*\*pylori\*\*\* \*\*\*urease\*\*\* to mucin and lipopolysaccharides  
contributes to gastric mucosal colonization by the bacterium based on in  
vivo targeting experiments using specific polysaccharides in a mouse model  
with acute infection. Acid-functional \*\*\*urease\*\*\*-homing  
polysaccharides that can interfere with \*\*\*urease\*\*\*-mucin or  
\*\*\*H\*\*\*. \*\*\*pylori\*\*\* whole cell-mucin interaction in vitro can  
significantly interfere with colonization by the bacterium in vivo.

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COST IN U.S. DOLLARS	SINCE FILE	TOTAL
	ENTRY	SESSION
FULL ESTIMATED COST	63.42	63.63
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
	ENTRY	SESSION
CA SUBSCRIBER PRICE	-0.69	-0.69

STN INTERNATIONAL LOGOFF AT 07:51:41 ON 20 JAN 2004